



California Medical Device Recall Information



Recall Name

**Bausch and Lomb Recalls 27G Sterile Cannula
Packed in Bausch and Lomb Amvisc 1.2% Sodium Hyaluronate and
Amvisc Plus 1.6% Sodium Hyaluronate Ophthalmic Viscosurgical Devices
Due to the Possibility of the Cannula Leaking or Detaching**

Recall Date	Product Description	Recalling Firm	Recall Reason
11/14/12	27G Sterile Cannula Component packed in two Bausch and Lomb products (see below)	Bausch and Lomb, Inc. Rochester, NY	<i>Cannula suspected of leaking or detaching from the syringe</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	Bausch and Lomb 27G Sterile Cannulas are found in: <ul style="list-style-type: none">• Amvisc 1.2% Sodium Hyaluronate (Models: 59051, 59081, 59051L, 59081L)• Amvisc Plus 1.6% Sodium Hyaluronate Ophthalmic Visco- surgical device (Models: 60081, 60051, 60051L, 60081L) List of Suspected Lots Recalled	CA , nationwide	Manufactured between December 2009 and August 2012

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm336413.htm>

